Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1-7. (Canceled)

- 8. (Currently Amended) A method of inhibiting angiogenesis in pathological conditions where increased angiogenesis and coincidental vascular perfusion are clinically detrimental, comprising the steps of: producing an AT₄ receptor ligand, having a structure selected from the group consisting of with the structure NH₃⁺-norleucine-tyrosine-isoleucine-histidine-COO, and norleucine-tyrosine-isoleucine-(6-amino-hexanoic acid) CONH₂; or norleucine-tyrosine-leucine-\frac{1}{2} (CH₂-HN₂)³⁻⁴-histidine-proline-phenylalanine) COO, and administering the AT₄ receptor ligand.
- 9. (Currently Amended) The method of inhibiting angiogenesis accordingly according to claim 8 or claim 29, further comprising the delivery of the AT₄ receptor ligand locally.
- 10. (Currently Amended) The method of inhibiting angiogenesis according to claim 8 or claim 29, further comprising the delivery of the AT₄ receptor ligand intravascularly.
- 11. (Currently Amended) The method of inhibiting angiogenesis according to claim 8 or claim 29, further comprising the delivery of the AT₄ receptor ligand intramuscularly.
- 12. (Currently Amended) The method of inhibiting angiogenesis according to claim 8 or claim 29, further comprising the delivery of the AT₄ receptor ligand intraperitoneally.

- 13. (Currently Amended) The method of inhibiting angiogenesis according to claim 8 or claim 29, further comprising the delivery of the AT₄ receptor ligand subcutaneously.
- 14. (Currently Amended) The method of inhibiting angiogenesis according to claim 8 or claim 29, further comprising the delivery of the AT₄ receptor ligand orally.
- of solid tumors, comprising the steps of: producing an AT₄ receptor ligand, having a structure selected from the group consisting of: with the structure NH₃⁺-norleucine-tyrosine-isoleucine-histidine-COO, and norleucine-tyrosine-isoleucine-(6-amino-hexanoic acid) CONH₂; or norleucine-tyrosine-leucine-\(\foat{V}\) (CH₂-HN₂)³⁻⁴-histidne-proline-phenylalanine) COO, and administering the AT₄ receptor ligand.
- 16. (Currently Amended) The method of inhibiting the growth and metastasis of solid tumors according to claim 15 or claim 30, further comprising delivery of the AT₄ receptor ligand locally.
- 17. (Currently Amended) The method of inhibiting the growth and metastasis of solid tumors according to claim 15 or claim 30, further comprising the delivery of the AT₄ receptor ligand intravascularly.
- 18. (Currently Amended) The method of inhibiting the growth and metastasis of solid tumors according to claim 15 or claim 30, further comprising the delivery of the AT₄ receptor ligand intramuscularly.
- 19. (Currently Amended) The method of inhibiting the growth and metastasis of solid tumors according to claim 15 or claim 30, further comprising the delivery of the AT₄ receptor ligand intraperitoneally.

- 20. (Currently Amended) The method of inhibiting the growth and metastasis of solid tumors according to claim 15 or claim 30, further comprising the step of applying the AT₄ receptor ligand subcutaneously.
- 21. (Currently Amended) The method of inhibiting the growth and metastasis of solid tumors according to claim 15 or claim 30, further comprising the step of applying the AT₄ receptor ligand orally.
- 22. (Currently Amended) A method of inhibiting the growth and metastasis of breast cancer, comprising the steps of: producing an AT₄ receptor ligand, having a structure selected from the group consisting of: with the structure NH₃⁺-norleucine-tyrosine-isoleucine-histidine-COO, and norleucine-tyrosine-isoleucine-(6-amino-hexanoic acid) CONH₂; or norleucine-tyrosine-leucine-Ψ-(CH₂-HN₂)³⁻⁴-histidne-proline-phenylalanine)-COO and administering the AT₄ receptor ligand.
- 23. (Currently Amended) The method of inhibiting the growth and metastasis of breast cancer according to claim 22 or claim 31, further comprising the delivery of the AT₄ receptor ligand locally to the tumor.
- 24. (Currently Amended) The method of inhibiting the growth and metastasis of breast cancer according to claim 22 or claim 31, further comprising the delivery of the AT₄ receptor ligand intravascularly.
- 25. (Currently Amended) The method of inhibiting the growth and metastasis of breast cancer according to claim 22 or claim 31, further comprising the delivery of the AT₄ receptor ligand intramuscularly.
- 26. (Currently Amended) The method of inhibiting the growth and metastasis of breast cancer according to claim 22 or claim 31, further comprising the delivery of the AT₄ receptor ligand intraperitoneally.

- 27. (Currently Amended) The method of inhibiting the growth and metastasis of breast cancer according to claim 22 or claim 31, further comprising the delivery of the AT₄ receptor ligand subcutaneously.
- 28. (Currently Amended) The method of inhibiting the growth and metastasis of breast cancer according to claim 22 or claim 31, further comprising the delivery of the AT₄ receptor ligand orally.
- 29. (New) A method of inhibiting angiogenesis in pathological conditions where increased angiogenesis and coincidental vascular perfusion are clinically detrimental, comprising the steps of: producing an AT₄ receptor ligand having a structure of norleucine-tyrosine-leucine-Ψ-(CH₂-NH₂)³⁻⁴-histidine-proline-phenylalanine-COO; and administering the AT₄ receptor ligand.
- 30. (New) A method of inhibiting the growth and metastasis of solid tumors, comprising the steps of: producing an AT₄ receptor ligand having a structure of: norleucine-tyrosine-leucine-Ψ-(CH₂-NH₂)³⁻⁴-histidine-proline-phenylalanine-COO; and administering the AT₄ receptor ligand.
- 31. (New) A method of inhibiting the growth and metastasis of breast cancer, comprising the steps of: producing an AT₄ receptor ligand having a structure of norleucine-tyrosine-leucine-Ψ-(CH₂-NH₂)³⁻⁴-histidine-proline-phenylalanine-COO; and administering the AT₄ receptor ligand.